

[54] METAL PERMANGANATE AND METAL PERIODATE ORGANIC ELECTROLYTE CELLS

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[52] U.S. Cl..... 136/6 LN; 136/100 R

[51] Int. Cl. 2..... H01M 35/02

[58] Field of Search 136/100, 6, 137, 138

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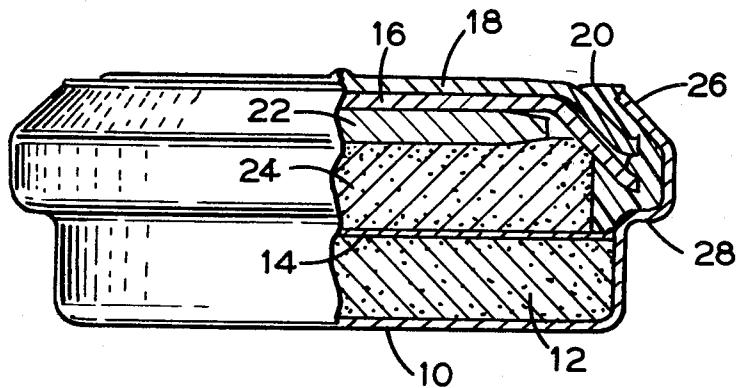
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[57] ABSTRACT

A cathodic active material is described utilizing the permanganates and periodates of heavy metals and preferably of silver. These materials are particularly suitable for use as positive electrodes in non-aqueous electrolyte cells and such electrodes and the cells are described. A method for fabrication of the active material into cathodes is included.

5 Claims, 7 Drawing Figures



SHEET 1 OF 3

FIG.1

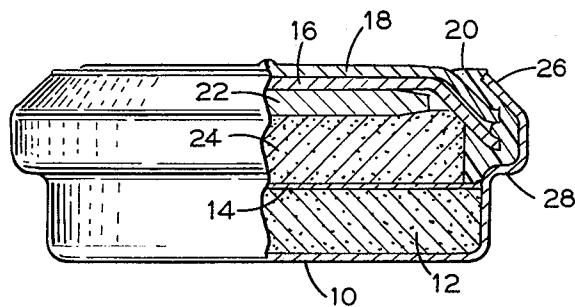


FIG.2

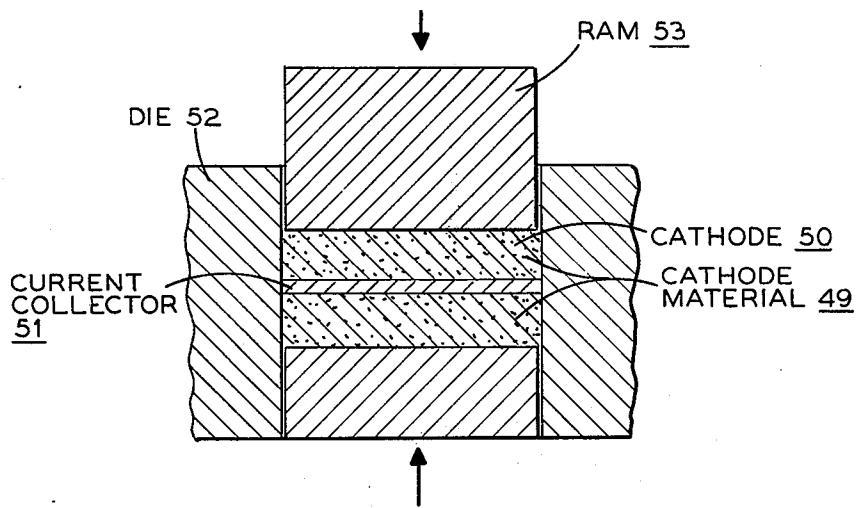
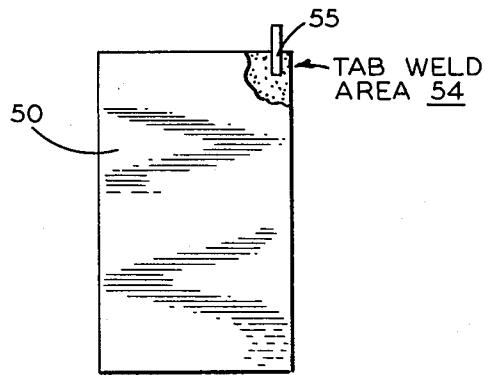


FIG.3



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FIG. 4

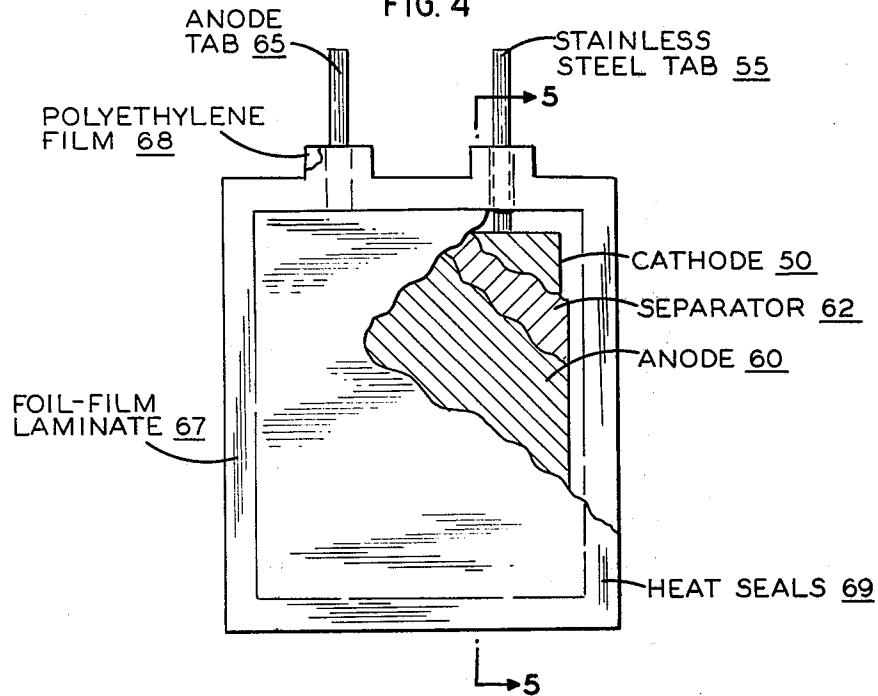
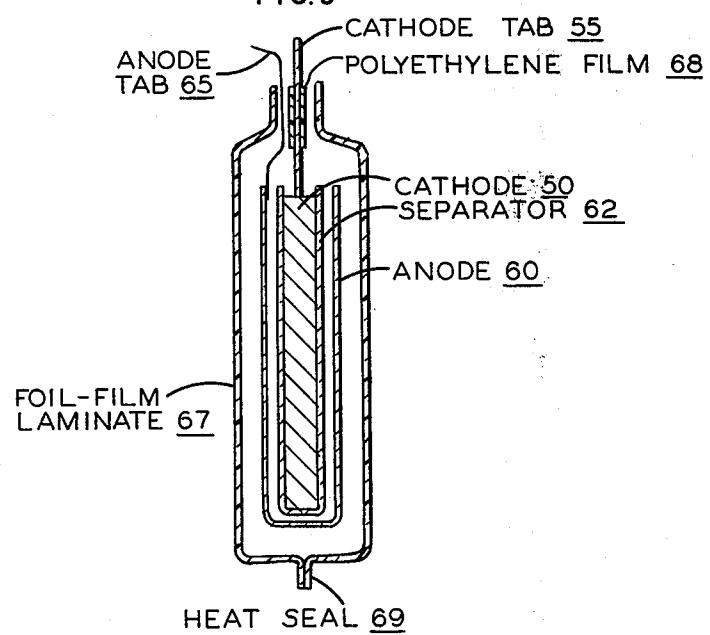


FIG. 5



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FIG. 6

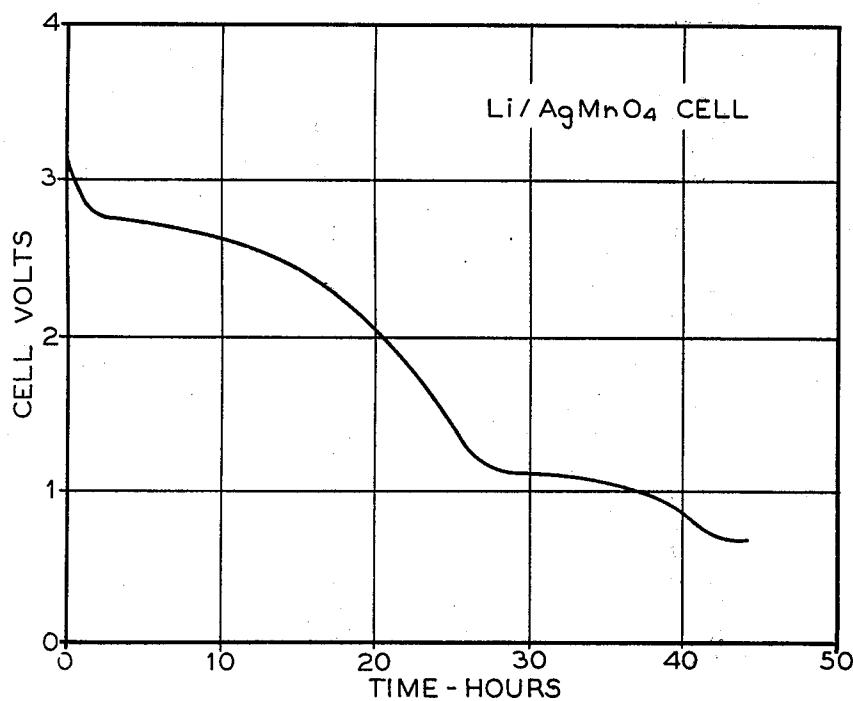
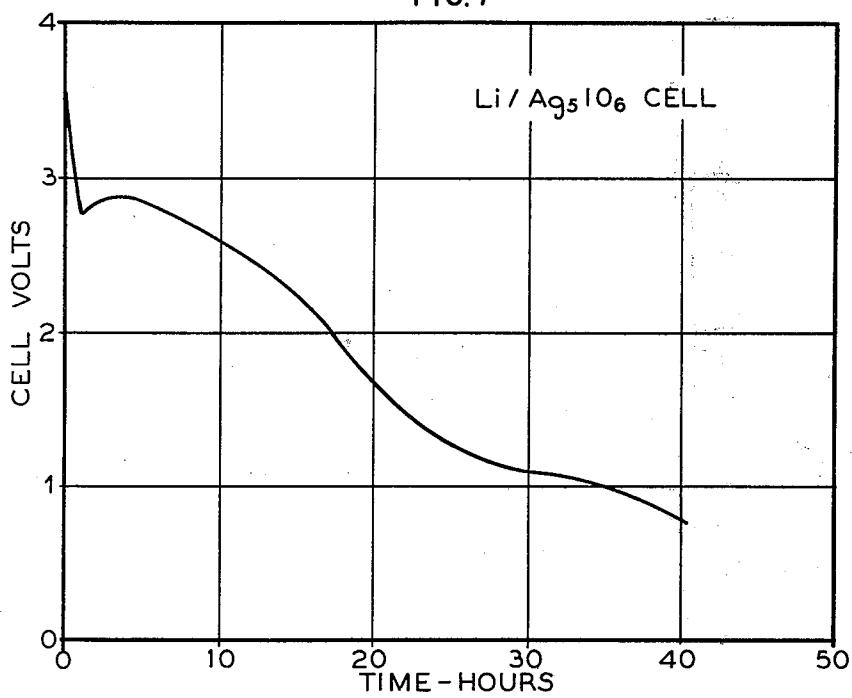


FIG. 7



**METAL PERMANGANATE AND METAL
PERIODATE ORGANIC ELECTROLYTE CELLS**

FIELD OF THE INVENTION

This invention relates to high-energy-density electrochemical energy generators and more particularly to such generating cells and batteries having organic electrolytes and utilizing as cathodic material the permanganates and periodates of certain metals.

BACKGROUND OF THE INVENTION

Metal permanganates and iodates are unstable in acid or alkaline solutions. For this reason it has not previously been possible to use such materials as depolarizers for conventional acid or alkaline primary or secondary cells.

The object of this invention is to provide novel electrode chemical energy generators with:

- a. high voltages,
- b. high energy densities,
- c. high material utilization efficiencies
- d. no spontaneous gasing

This invention is based upon the use, as active cathode material of the oxy acids of manganese and iodine in the form of their metal salts with certain heavy metals. The permanganates and periodates of copper, silver, iron, cobalt, and mixtures thereof are particularly suitable as depolarizers for this invention. Such active cathodic materials are used in conjunction with non-aqueous electrolytes in electrochemical generators. The electrochemical generators for the utilization of such active materials include, as anodic material, active light metals preferably of the type of lithium, sodium, potassium, etc.

Within the ambit of this invention are the novel active cathodic materials; cathodes prepared from such novel cathodic material; methods of making such cathodes from said active materials and the complete electrochemical generators using such active cathodic and anodic materials in conjunction with non-aqueous electrolytes. Broadly stated according to the principles of the present invention, there is provided an electric cell comprising a light metal anode, a heavy metal permanganate or periodate cathode, and an organic electrolyte.

Light metals suitable for the purpose of the present invention are selected from among lithium, sodium, potassium, beryllium, calcium, magnesium, and aluminum. In some cases, it is desirable to amalgamate the surface of the anode metal with mercury as the performance of certain light metal anodes for such cells can be considerably improved by such amalgamation. By surface alloying these metals with mercury by chemical displacement of the latter from solutions of a mercury salt, a notable improvement of the anodes is provided. The improvement is particularly striking with anodes formed of aluminum, magnesium, or their alloys. Amalgamation may be successfully carried out by bringing the anode metal into contact with mercuric salts dissolved in N:N dimethyl formamide, gamma-butyrolactone, acetone or in other suitable solvents. Among useful mercuric salts for such solution are mercuric chloride, mercuric nitrate, mercuric acetate, mercuric chlorate, mercuric bromide and mercuric thiocyanate. Generally speaking the principles of selection of a suitable system (solute-solvent) for amalgamation of the following:

- a. The mercury salt should be soluble in the solvent.
- b. The anion of the mercury salt should form salts with the anode metal, that are also soluble in the solvent employed to dissolve the mercury salt.

- c. The solvent should be stable in the presence of the anode metal and its amalgam, more particularly it should not decompose evolving hydrogen, as would be the case with acetic acid.

The active metal anodes are amalgamated to reduce local action on the anode surface thus providing consistent and reproducible electrochemical results. Specifically it has been noted, with aluminum and magnesium anodes, that amalgamation assures elimination of the usual time lag for initiation of the current flow. With anodes of such metals and their alloys the usual time lag is reduced to less than 10^{-3} seconds.

The novel cathodic material or depolarizers used in the present invention are the heavy metal salts of the oxy acids of manganese and iodine. The permanganates and periodates (MnO_4^{-1}) or (IO_6^{-5}) of copper, silver, iron, cobalt, nickel, mercury, thallium, lead, and bismuth and mixtures thereof are particularly suitable as the active cathodic or depolarizer materials for cells to be used in conjunction with the above mentioned active light metal anodes. The silver permanganates, $AgMnO_4$ and the silver periodates, Ag_5IO_6 are preferred.

Suitable electrolytes may be made by dissolving organic or inorganic salts of light metals in the organic solvents. For example, 1-2 Molar solutions of lithium perchlorate, or lithium aluminum chloride dissolved in tetrahydrofuran solvent constitute a suitable organic electrolyte. Other light metal salts are based upon the cations of the light metals such as lithium, sodium, beryllium, calcium, magnesium or aluminum with anions such as perchlorate, tetrachloroaluminate, tetrafluoroborate, chloride, hexafluorophosphate, hexafluoroarsenates, etc. dissolved in organic solvents. Among suitable organic solvents are tetrahydrofuran, propylene carbonate, dimethyl sulfide, dimethyl sulfoxide, N-nitrosodimethylamine, gamma-butyrolactone, dimethyl carbonate, methyl formate, butyl formate, acetonitrile and N:N dimethyl formamide.

This invention in addition to the electrochemical system mentioned above also contemplates cells and batteries utilizing this system which are simple in construction reliable and safe in operation, and which may be readily manufactured and sold on a practical and commercial scale at a low cost. Similarly included within the ambit of this invention is the method of manufacture of the cathodes from the cathodic materials. These and other aspects and objects and advantages of the present invention will be, apparent from the following description taken in conjunction with the accompanying drawing in which:

FIG. 1 is a vertical sectional view, having parts in elevation embodying a commercial cell according to this invention.

FIG. 2 shows a method of manufacture of the cathode according to one aspect of this invention ready for compression in a molding die.

FIG. 3 shows a finished cathode according to one aspect of this invention including a tab welded area for affixing current collecting leads. The current collectors

of FIG. 4 shows a side view of a cell in its individually sealed envelope useful according to another aspect of this invention. FIG. 5 shows a sectional view of the cell according to FIG. 4, said section being taken along line

5—5 of FIG. 4. FIG. 6 is a discharge curve of a lithium-
/AgMnO₄ cell according to this invention.

FIG. 7 is a discharge curve of a lithium/Ag₅IO₆ cell
according to this invention.

Referring now particularly to FIG. 1 of the drawing,
the structure used and the procedure followed in as-
sembling a practical cell and embodying this one aspect
of this invention will be described.

Reference numeral 10 denotes a cathode container
made of stainless steel, in which there is compressed a
body of cathode mix 12. This cathode mix employing
the cathodic material of this invention is a pre-mixed
and pre-slugged mixture of the powdered depolarizer
comprising the heavy metal permanganates or perio-
dates and graphite or any similar conductive carbon in
a weight ratio of approximately seven parts of depolar-
izer to three parts of graphite. The pre-mixture is
bonded with approximately 3% by weight of a aqueous
dispersion of polytetrafluoroethylene commercially
available as colloidal Teflon (Dupont de Nemours Co.
Inc.). After the water in the dispersion has been dis-
placed by an organic solvent such as isopropanol or
benzene, the cathodic mixture is compressed. After
compression, it is preferred to dry and cure the previ-
ously compressed cathode mixture at temperatures of
approximately 300°C for about 2 hours. A thin barrier
or separator layer 14 of a microporous inert material
such as fiber glass is then placed in the cathode con-
tainer 10 on top of the compressed cathode mix 12.

The top closure of the cell comprises inner top disc
16 made of stainless steel and outer top disc 18, like-
wise made of stainless steel, the two top discs having
their center portions nested in and their edge portions
slightly separated from each other. An insulating and
sealing grommet 20 of a suitable elastomer such as neo-
prene or polyethylene, is stretched or molded around
the circumferential edges of top discs 16 and 18. A de-
tailed description of the assembly and operation of this
so called "double top" structure will be found in Wil-
liams U.S. Pat. No. 2,712,565 and Clune U.S. Pat. No.
3,096,216 to which reference is hereby made. In inner
top disc 16 there is anode 22 cut from a plate of the
anode metal, for example lithium, to fit the inner di-
mension of said disc. A body of an electrolyte absor-
bent 24, such as cotton is interposed between anode 22
and the cathode mix 12. Onto this electrolyte absor-
bent 24 is deposited about 0.5 cc's of an electrolyte such
as 1 molar lithium perchlorate dissolved in propylene
carbonate. Thereupon, the cell is assembled and per-
manently sealed by crimping bown enlarged mouth
portions 26 of cathode container 10 over grommet 20
which is thus strongly compressed between the said
mounth portions and the annular shoulder 28 of the con-
tainer. Any excess electrolyte present is squeezed out
during the final assembly of the cell.

FIGS. 2 and 3 show a schematic method for manufac-
ture of cathodes useful in the manufacture of sealed in-
dividual cells which may be assembled into batteries. In
the cathode construction according to this aspect of the
invention, the cathodes 50 are fabricated by compris-
ing a pre-mixed and pre-slugged mixture 49 of pow-
dered depolarizer and graphite in a 7:3 weight ratio
with approximately three parts by weight of a binder.
The preferred binder is 3% by weight of a aqueous dis-
persion of polytetrafluoroethylene, commonly known
as colloidal Teflon (Dupont de Nemours Co. Inc.) The
water of this aqueous dispersion is then displaced by
adding a sufficient amount of an organic solvent such

as isopropanol, benzene, etc., to form a paste added to
the mix. The paste is then thoroughly mixed to form a
easily pliable dough. Any excess solvent is decanted.
The cathodes 50 are molded on, and around nickel cur-
rent collector 51, preferably fabricated from metal
screening or expanded metal, by placing layers of the
dough above and below the current collector 51 which
is in turn placed in a rectangular die 52 and then press-
ing the dough at pressures 70-80,000 lbs/in² by ram 53.
The process is shown schematically in FIG. 2. The ex-
cess solvent such as isopropanol is squeezed out of the
dough and there results a compact rectangular cathode
50 with adequate mechanical integrity for further treat-
ment. This cathode 50 is then dried in air and prefer-
ably cured at a temperature between 200°-350° from 1½
to 3 hours. Optimally the curing treatment is 300°C for
2 hours. The curing process enhances the mechanical
integrity of the cathode even further. The electrical
conductivity of such a cathode is more than adequate
for its purpose. In the above method, the ratio of active
material to graphite or other conductive inert additive
and the ratio of the total mix to binder can be varied
widely.

25 The nature of the binder can also be varied to a con-
siderable extent. The binder may include both organic
and inorganic organic compounds. Organic binders in
addition to the colloidal Teflon are polyethylene dis-
solved in xylene, ethyl cellulose dissolved in xylene,
30 and so forth. Other polyolefin polymers may similarly
be used. Inorganic binders which may be used include
plaster of Paris proprietary dental cements. The pur-
pose of the pressure is to mechanically consolidate the
35 cathode mixture. The preferred pressure is 70,000
lbs/in².

When curing the cathodes, the curing temperature
can be varied from 100° to 400°C and the curing time
should be varied according to the temperature.

40 A corner 54 of the cathode 50 after heat treatment
made according to the above procedure, is scraped to
bare a portion of the current collector 51 to which a tab
55 is spot welded for electrical connection. The fin-
ished cathode is shown in FIG. 3. This cathode, used in
45 conjunction with lithium anodes is shown in a side view
of the cell at FIG. 4 including partial cut-outs.

A section of FIG. 4 long lines 5—5 is shown in FIG.
5. The cathode 50 is used in conjunction with two lith-
ium anodes 60, prepared by pressing two rectangular
50 pieces of lithium metal approximately 0.02 inches
thick, on expanded stainless steel. This cell is con-
structed in parallel plate configuration. A layer of filter
paper 62 is interposed between the lithium anode 60
and the cathode 50 to provide a separator 62 and an
55 electrolyte absorber 62. The cell is enclosed in a poly-
mer and foil-laminate enclosure 67. These foil-laminate
film cells are heat-sealed around the periphery prefer-
ably by heat seals 69. At the upper end where the cath-
ode tabs 55 and anode tabs 65 leave the cell structure
60 proper, portions of polyethylene film 68 are affixed to
the tabs to provide leak proof seals. Before sealing the
cells 3cc's of 1.0 molar solution of lithium perchlorate
(LiClO₄) in tetrahydrofuran is added to each cell en-
closure and the cell is permanently heat sealed. The
65 cells according to this type of construction are cathode
limited, i.e., the cathode has a lower capacity than the
anode. They may be assembled into batteries by series
or parallel connections.

OPERATION

All the cells are discharged using a constant current of 4.5 ma corresponding to a current density of 1 ma/cm² based on the geometric area of both the sides of the cathode. This corresponds to a 10-30 hour rate. The operating characteristics of these cells under the above type of load is shown in Table 1.

(a) comprising a cathodic material selected from the group consisting of the permanganates and periodates of heavy metals from the group consisting of copper, silver, iron, cobalt, nickel, mercury, thallium, lead, and bismuth, and mixtures thereof; said negative electrode
 5 (b) comprising an active light metal selected from the group consisting of lithium, sodium, potassium, beryllium, calcium, magnesium, and aluminum and alloys

TABLE 1

Operating Characteristics of Various Li/Metal Permanganate and Li/Metal Periodate Organic Electrolyte Cells				
Example No.	Cell	Open Circuit Voltage	Average Operating Voltage	Cathode Material Utilization Efficiency (%)
1.	Li/AgMnO ₄ (FIG. 6)	3.5	2.4	70%
2.	Li/Ag ₃ IO ₆ (FIG. 7)	3.5	2.3	100%
	Both of the cells showed high open circuit and operating voltages compared to the conventional alkaline cells. The discharge characteristics of these cells are shown in FIGS. 6 and 7. The systems Li/AgMnO ₄ , Li/Ag ₃ IO ₆ , are particularly important for their high and exceedingly steady output voltage. The assumed cathode reactions for all these cells given in Table II below:			
Ex. 1.	AgMnO ₄ + 4Li ⁺ + 4e → Ag + MnO ₂ + 2Li ₂ O			
Ex. 2.	Ag ₃ IO ₆ + 11Li ⁺ + 11e → 5Ag + LiIO + 5Li ₂ O The anodic reaction in all the above cells is Li → Li ⁺ + e.			

For the lithium anodes there may be substituted, in the same molar proportion sodium, potassium, rubidium, and cesium. Stoichiometric equivalents of beryllium, calcium, magnesium, barium, strontium and aluminum may be substituted for the lithium.

The present invention has been disclosed by reference to preferred embodiments thereof. However, variations and modifications may be resorted to by those skilled in the art without departing from the principles of this invention. All the variations and modifications are considered to be within the true spirit and scope of the present invention as disclosed in the foregoing description and including present art recognized equivalents for the various components.

What is claimed is:

1. An electrochemical energy generating cell comprising (a) at least one positive electrode; (b) at least one negative electrode; (c) an organic electrolyte with said electrodes disposed therein; said positive electrode

thereof; and said organic electrolyte (c) comprising a polar organic solvent having dissolved therein at least one ion-forming salt of an active light metal.

2. The cell according to claim 1 wherein said polar solvent is selected from the group of tetrahydrofuran, 30 N-nitrosodimethylamine, dimethyl sulfite, propylene carbonate, dimethyl sulfoxide, gamma-butyrolactone, dimethyl carbonate, methyl formate, butyl formate, acetonitrile and N:N dimethyl formamide.

3. The cell according to claim 2, wherein the salt dissolved in said polar organic solvent is selected from the group consisting of active light metal salts of perchlorates, tetrachloroaluminates, tetrafluoroborates, chlorides, hexafluorophosphates, and hexafluoroarsenates.

35 4. The cell according to claim 1, wherein the cathodic active material is silver permanganate.

5. The cell according to claim 1, wherein the cathodic active material is silver periodate.

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